



UNITED STATES PATENT AND TRADEMARK OFFICE

UNITED STATES DEPARTMENT OF COMMERCE
United States Patent and Trademark Office
Address: COMMISSIONER FOR PATENTS
P.O. Box 1450
Alexandria, Virginia 22313-1450
www.uspto.gov

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/936,759	11/07/2001	Richard A. Jefferson	076518-0150	8995
39124 : 7590 02/16/2005 CAROL NOTTENBURG 814 32ND AVE 5 SEATTLE, WA 98144			EXAMINER VOGEL, NANCY S	
			ART UNIT 1636	PAPER NUMBER

DATE MAILED: 02/16/2005

Please find below and/or attached an Office communication concerning this application or proceeding.

Office Action Summary	Application No.		Applicant(s)	
	09/936,759		JEFFERSON ET AL.	
	Examiner		Art Unit	
	Nancy T. Vogel		1636	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☐ Responsive to communication(s) filed on ____.
- 2a) ☐ This action is **FINAL**. 2b) ☒ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 36,37 and 60-63 is/are pending in the application.
- 4a) Of the above claim(s) ____ is/are withdrawn from consideration.
- 5) ☐ Claim(s) ____ is/are allowed.
- 6) ☒ Claim(s) 36,37 and 60-63 is/are rejected.
- 7) ☐ Claim(s) ____ is/are objected to.
- 8) ☐ Claim(s) ____ are subject to restriction and/or election requirement.

Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☒ The drawing(s) filed on ____ is/are: a) ☐ accepted or b) ☒ objected to by the Examiner.
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☒ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) ☒ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some * c) ☐ None of:
1. ☐ Certified copies of the priority documents have been received.
2. ☐ Certified copies of the priority documents have been received in Application No. ____.
3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).
- * See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- | | |
|---|---|
| 1) <input checked="" type="checkbox"/> Notice of References Cited (PTO-892) | 4) <input type="checkbox"/> Interview Summary (PTO-413)
Paper No(s)/Mail Date. ____. |
| 2) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948) | 5) <input type="checkbox"/> Notice of Informal Patent Application (PTO-152) |
| 3) <input checked="" type="checkbox"/> Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08)
Paper No(s)/Mail Date <u>9/12/02, 4/26/02</u> . | 6) <input checked="" type="checkbox"/> Other: <u>seq. alignment</u> . |

DETAILED ACTION

Claims 36, 37, and 60-63 are pending in the case.

Information Disclosure Statement

Receipt of Information disclosure statements on 9/12/02 and 4/26/02 is acknowledged. Certain citations have not been considered, since copies have not been received. These citations have been struck through and the information referred to therein has not been considered.

Drawings

New corrected drawings in compliance with 37 CFR 1.121(d) are required in this application because the text of some of the drawings cannot be read (Figures 5 and 5c). Applicant is advised to employ the services of a competent patent draftsman outside the Office, as the U.S. Patent and Trademark Office no longer prepares new drawings. The corrected drawings are required in reply to the Office action to avoid abandonment of the application. The requirement for corrected drawings will not be held in abeyance.

Oath/Declaration

The oath or declaration is defective. A new oath or declaration in compliance with 37 CFR 1.67(a) identifying this application by application number and filing date is required. See MPEP §§ 602.01 and 602.02.

The oath or declaration is defective because:

Art Unit: 1636

It improperly lists a US patent application, 09/270,957, as a foreign application to which applicants claim benefit under 35 USC 119 (a-d).

Priority

It is noted that this application appears to claim subject matter disclosed in prior Application No. 09/270,957, filed 3/17/99. A reference to the prior application must be inserted as the first sentence of the specification of this application or in an application data sheet (37 CFR 1.76), if applicant intends to rely on the filing date of the prior application under 35 U.S.C. 119(e) or 120. See 37 CFR 1.78(a). For benefit claims under 35 U.S.C. 120, the reference must include the relationship (i.e., continuation, divisional, or continuation-in-part) of all nonprovisional applications. Also, the current status of all nonprovisional parent applications referenced should be included.

If the application is a utility or plant application filed under 35 U.S.C. 111(a) on or after November 29, 2000, the specific reference to the prior application must be submitted during the pendency of the application and within the later of four months from the actual filing date of the application or sixteen months from the filing date of the prior application. If the application is a utility or plant application which entered the national stage from an international application filed on or after November 29, 2000, after compliance with 35 U.S.C. 371, the specific reference must be submitted during the pendency of the application and within the later of four months from the date on which the national stage commenced under 35 U.S.C. 371(b) or (f) or sixteen months from the filing date of the prior application. See 37 CFR 1.78(a)(2)(ii) and (a)(5)(ii). This

time period is not extendable and a failure to submit the reference required by 35 U.S.C. 119(e) and/or 120, where applicable, within this time period is considered a waiver of any benefit of such prior application(s) under 35 U.S.C. 119(e), 120, 121 and 365(c). A priority claim filed after the required time period may be accepted if it is accompanied by a grantable petition to accept an unintentionally delayed claim for priority under 35 U.S.C. 119(e), 120, 121 and 365(c). The petition must be accompanied by (1) the reference required by 35 U.S.C. 120 or 119(e) and 37 CFR 1.78(a)(2) or (a)(5) to the prior application (unless previously submitted), (2) a surcharge under 37 CFR 1.17(t), and (3) a statement that the entire delay between the date the claim was due under 37 CFR 1.78(a)(2) or (a)(5) and the date the claim was filed was unintentional. The Director may require additional information where there is a question whether the delay was unintentional. The petition should be addressed to: Mail Stop Petition, Commissioner for Patents, P.O. Box 1450, Alexandria, Virginia 22313-1450.

Specification

The amendment filed 4/22/02 is objected to under 35 U.S.C. 132 because it introduces new matter into the disclosure. 35 U.S.C. 132 states that no amendment shall introduce new matter into the disclosure of the invention. The added material which is not supported by the original disclosure is as follows: the amendments to 49 which changes nucleotide sequences; a nucleotide sequence on page 56.

Applicant is required to cancel the new matter in the reply to this Office Action.

Double Patenting

A rejection based on double patenting of the "same invention" type finds its support in the language of 35 U.S.C. 101 which states that "whoever invents or discovers any new and useful process ... may obtain a patent therefor ..." (Emphasis added). Thus, the term "same invention," in this context, means an invention drawn to identical subject matter. See *Miller v. Eagle Mfg. Co.*, 151 U.S. 186 (1894); *In re Ockert*, 245 F.2d 467, 114 USPQ 330 (CCPA 1957); and *In re Vogel*, 422 F.2d 438, 164 USPQ 619 (CCPA 1970).

A statutory type (35 U.S.C. 101) double patenting rejection can be overcome by canceling or amending the conflicting claims so they are no longer coextensive in scope. The filing of a terminal disclaimer cannot overcome a double patenting rejection based upon 35 U.S.C. 101.

Claims 36 is provisionally rejected under 35 U.S.C. 101 as claiming the same invention as that of claim 36 of copending Application No. 10/364,649. This is a provisional double patenting rejection since the conflicting claim has not in fact been patented.

The nonstatutory double patenting rejection is based on a judicially created doctrine grounded in public policy (a policy reflected in the statute) so as to prevent the unjustified or improper timewise extension of the "right to exclude" granted by a patent and to prevent possible harassment by multiple assignees. See *In re Goodman*, 11 F.3d 1046, 29 USPQ2d 2010 (Fed. Cir. 1993); *In re Longi*, 759 F.2d 887, 225 USPQ 645 (Fed. Cir. 1985); *In re Van Ornum*, 686 F.2d 937, 214 USPQ 761 (CCPA 1982); *In re Vogel*, 422 F.2d 438, 164 USPQ 619 (CCPA 1970); and, *In re Thorington*, 418 F.2d 528, 163 USPQ 644 (CCPA 1969).

A timely filed terminal disclaimer in compliance with 37 CFR 1.321(c) may be used to overcome an actual or provisional rejection based on a nonstatutory double patenting ground provided the conflicting application or patent is shown to be commonly owned with this application. See 37 CFR 1.130(b).

Effective January 1, 1994, a registered attorney or agent of record may sign a terminal disclaimer. A terminal disclaimer signed by the assignee must fully comply with 37 CFR 3.73(b).

Claim 37 is provisionally rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claim 36 of copending Application No. 10/364,649. Although the conflicting claims are not identical, they are not patentably distinct from each other because the instant claim 37 recites beta glucuronidase which is encompassed by the beta-glucuronidases recited in copending 10/364,649, claim 36, since an isolated beta glucuronidase having 90% identity with the sequence shown in SEQ ID NO:6, would be encoded by a nucleic acid molecule that would hybridize under stringent conditions to the complement of nucleotides 1-1689 of SEQ ID NO:14.

This is a provisional obviousness-type double patenting rejection because the conflicting claims have not in fact been patented.

Claims 60-63 are provisionally rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claim 36 of copending Application No. 10/364,649 in view of Hochuli et al. (Bio/Technology 6:1321-1325, 1988)

Claim 36 of copending application 10/364,649 teaches an isolated beta-glucuronidase encoded by a nucleic acid molecule comprising nucleotides 1-1689 of Figure 4I-J (SEQ ID NO:14) or by a nucleic acid molecule that hybridizes under stringent conditions to the complement of said nucleic acid molecule. This nucleic acid molecule, and the beta-glucuronidase encoded by it, is the same as that disclosed in

Fig. 4I-J, or SEQ ID NO:14 of the instant application (nucleic acid) and SEQ ID NO:6 (protein).

The difference between the copending application and the instant claims is that the instant claims recite a fusion protein comprising the isolated beta-glucuronidase and a peptide, which may be hexa-His.

However, Hochuli et al. disclose protein fusions between a protein of interest, and a peptide of six histidine residues, i.e. "hexa-His", and the usefulness of said fusion protein for purification of the protein of interest (see . It would have been obvious to those of ordinary skill in the art, to have modified the protein which is the beta-glucuronidase of SEQ ID NO:14 in the US patent application 10/364,649, by fusing it to a peptide such as hexa-His, as taught by Hochuli et al., since the references generally concern the production and isolation of proteins of interest using genetic techniques, and since Hochuli et al. teach general techniques that are known in the art to be useful for isolation of proteins of interest or for bioassays. One would have been motivated to do so by the desire to obtain in purified form, the protein of interest which is disclosed by the US patent application, 10/364,649, since it is well known in the art that purified products have the advantage of lack of contaminants. Based upon the teachings of the cited references, the high skill of one of ordinary skill in the art, and absent evidence to the contrary, there would have been a reasonable expectation of success to result in the claimed invention.

This is a provisional obviousness-type double patenting rejection.

Claims 60-63 are provisionally rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claim 36 of copending Application No. 10/364,649 in view of Diamandis et al. (Clin. Chem. 37, 625, 1991) (cited by applicants).

Claim 36 of copending application 10/364,649 teaches an isolated beta-glucuronidase encoded by a nucleic acid molecule comprising nucleotides 1-1689 of Figure 4I-J (SEQ ID NO:14) or by a nucleic acid molecule that hybridizes under stringent conditions to the complement of said nucleic acid molecule. This nucleic acid molecule, and the beta-glucuronidase encoded by it, is the same as that disclosed in Fig. 4I-J, or SEQ ID NO:14 of the instant application (nucleic acid) and SEQ ID NO:6 (protein).

The difference between the copending application and the instant claims is that the instant claims recite a fusion protein comprising the isolated beta-glucuronidase and a peptide, which may be streptavidin.

However, Diamandis et al. teach fusion proteins between an enzyme of interest and streptavidin, and the usefulness of said fusion protein for such applications as immunoassays, flow cytometry, cell sorting, and Western blots (see Table 2 and pages 631-634). It would have been obvious to those of ordinary skill in the art, to have modified the protein which is the beta-glucuronidase of SEQ ID NO:14 in the US patent application 10/364,649, by fusing it to a peptide such as streptavidin, as taught by Diamandis et al., since the references concern enzymes of interest, and since Diamandis et al. teach general techniques of making enzyme- streptavidin fusions that

Art Unit: 1636

are known in the art to be applicable to enzymes whose activity is known, for use in bioassays. One would have been motivated to do so by the desire to obtain a beta-glucuronidase fusion useful for assays such as immunoassays. Based upon the teachings of the cited references, the high skill of one of ordinary skill in the art, and absent evidence to the contrary, there would have been a reasonable expectation of success to result in the claimed invention.

This is a provisional obviousness-type double patenting rejection.

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

Claims 36, 37, and 60 are rejected under 35 U.S.C. 103(a) as being unpatentable over Nelson et al. (Nature, 399, 323-3329, 1999) in view of Jefferson et al. (GB 2 197 653, cited by applicants).

Nelson et al. disclose a beta-glucuronidase from *Thermatoga maritima*, which is encoded by a nucleic acid molecule comprising nucleotides 1-1689 of SEQ ID NO:14 of the instant application, or which comprises the amino acid sequence of SEQ ID NO:6 of the instant application (see attached sequence alignments). The difference between the reference and the instant application is that the beta-glucuronidase is isolated.

However, Jefferson et al. disclose the isolation of beta-glucuronidase from a bacteria (see pages 12, 16-17). It would have been obvious to one of ordinary skill in the art, to have isolated the beta-glucuronidase which is disclosed by Nelson et al. using such well known techniques as that disclosed by Jefferson for the isolation of beta-glucuronidase, since both references disclose beta-glucuronidases from bacteria and their sequences. One would have been motivated to do so by the well known usefulness of purified or isolated proteins, which are free of contaminants and thus in a useful form.

Claims 36, 37, 60-63 are rejected under 35 U.S.C. 103(a) as being unpatentable over Nelson et al. in view of Jefferson et al. as applied to claims 36 and 37 above, and further in view of Hochuli et al. (Bio/Technology 6:1321-1325, 1988).

Nelson et al. and Jefferson et al. are cited for the reasons set forth above.

The difference between the references and the instant claims is that the instant claims recite a fusion protein comprising the isolated beta-glucuronidase and a peptide, which may be hexa-His.

However, Hochuli et al. disclose protein fusions between a protein of interest, and a peptide of six histidine residues, i.e. "hexa-His", and the usefulness of said fusion protein for purification of the protein of interest (see . It would have been obvious to those of ordinary skill in the art, to have modified the protein which is the beta-glucuronidase taught by Nelson et al and Jefferson, by fusing it to a peptide such as hexa-His, as taught by Hochuli et al., since the references generally concern the

Art Unit: 1636

production and isolation of proteins of interest using genetic techniques, and since Hochuli et al. teach general techniques that are known in the art to be useful for isolation of proteins of interest or for bioassays. One would have been motivated to do so by the desire to obtain in purified form, the protein of interest which is disclosed by the US patent application, 10/364,649, since it is well known in the art that purified products have the advantage of lack of contaminants. Based upon the teachings of the cited references, the high skill of one of ordinary skill in the art, and absent evidence to the contrary, there would have been a reasonable expectation of success to result in the claimed invention.

Claims 36, 37, 60-63 are rejected under 35 U.S.C. 103(a) as being unpatentable over Nelson et al. in view of Jefferson et al. as applied to claims 36 and 37 above, and further in view of Diamandis et al. (Clin. Chem. 37, 625, 1991) (cited by applicants).

Nelson et al. and Jefferson et al. are cited for the reasons set forth above.

The difference between the copending application and the instant claims is that the instant claims recite a fusion protein comprising the isolated beta-glucuronidase and a peptide, which may be streptavidin.

However, Diamandis et al. teach fusion proteins between an enzyme of interest and streptavidin, and the usefulness of said fusion protein for such applications as immunoassays, flow cytometry, cell sorting, and Western blots (see Table 2 and pages 631-634). It would have been obvious to those of ordinary skill in the art, to have modified the protein which is the beta-glucuronidase as taught by Nelson et al. and

Art Unit: 1636

Jefferson, by fusing it to a peptide such as streptavidin, as taught by Diamandis et al., since the references concern enzymes of interest, and since Diamandis et al. teach general techniques of making enzyme- streptavidin fusions that are known in the art to be applicable to enzymes whose activity is known, for use in bioassays. One would have been motivated to do so by the desire to obtain a beta-glucuronidase fusion useful for assays such as immunoassays. Based upon the teachings of the cited references, the high skill of one of ordinary skill in the art, and absent evidence to the contrary, there would have been a reasonable expectation of success to result in the claimed invention.

Conclusion

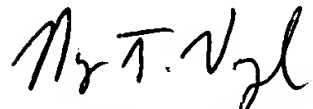
No claims are allowed.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Nancy T. Vogel whose telephone number is (571) 272-0780. The examiner can normally be reached on 7:00 - 3:30, Monday - Friday.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Irem Yucel, Ph.D. can be reached on (571) 272-0781. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Art Unit: 1636

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).



Nancy T. Vogel, Ph.D.
Patent Examiner

Db 61 ELSQKHIRLYFAAVNTDCEVFLNGEKVGENHIEYLPFEVDVTGKXSGENELRVVENRL 120
QY 121 KVGGFPSKVPDSCGHTVGFSGSPPPANFDFPYGGIIRPVLIEFTDHARILDIMWDTSES 180
Db 121 KVGGFPSKVPDSCGHTVGFSGSPPPANFDFPYGGIIRPVLIEFTDHARILDIMWDTSES 180
QY 181 EPEKKLGKVKKIEVSEAVGQEMTIKLGEEKKIRTSNRFVEGEFLENARFWSLEDPY 240
Db 181 EPEKKLGKVKKIEVSEAVGQEMTIKLGEEKKIRTSNRFVEGEFLENARFWSLEDPY 240
QY 241 LYPKLEKDEYTLDIGIRTI SWDEKRLYLNGKPVFLKGFGKHEEPVLGGTFYPLMI 300
Db 241 LYPKLEKDEYTLDIGIRTI SWDEKRLYLNGKPVFLKGFGKHEEPVLGGTFYPLMI 300
QY 301 KDFNLKWINANSFRTSHYPYSEEWDLADRLGILVIDEAPHVGI TRHYNPETOKIAED 360
Db 301 KDFNLKWINANSFRTSHYPYSEEWDLADRLGILVIDEAPHVGI TRHYNPETOKIAED 360
QY 361 NIRMIDRHKHPSVIMSVANEPESNHPDAEGFFKALYETANENDRTRPVVMVSMMDAP 420
Db 361 NIRMIDRHKHPSVIMSVANEPESNHPDAEGFFKALYETANENDRTRPVVMVSMMDAP 420
QY 421 DERTRDVALKYPDIVCVNRYGWTYYQRIEEGLQALEKDIEELYARHRKPIFVTEFGAD 480
Db 421 DERTRDVALKYPDIVCVNRYGWTYYQRIEEGLQALEKDIEELYARHRKPIFVTEFGAD 480
QY 481 AIAGIHYDPQPMFSEEQAEIYEKTI RLLKKDYII GTHVMAFADFKTPQNVRRPILNHK 540
Db 481 AIAGIHYDPQPMFSEEQAEIYEKTI RLLKKDYII GTHVMAFADFKTPQNVRRPILNHK 540
QY 541 GVFTDRDQPKLVAVHLRLMSEV 563
Db 541 GVFTDRDQPKLVAVHLRLMSEV 563

RESULT 2
Q97UI1 PRELIMINARY; PRT; 570 AA.

ID Q97UI1 AC Q97UI1; PRT; 570 AA.
AC Q97UI1; PRT; 570 AA.
DT 01-OCT-2001 (TReMBLrel. 18, Created)
DT 01-OCT-2001 (TReMBLrel. 18, Last sequence update)
DT 01-OCT-2003 (TReMBLrel. 25, Last annotation update)
DE Beta-glucuronidase (Gusb) (EC 3.2.1.31).
GN Name=gusb; Ordered locusNames=SCO3036;
OS Sulfolobus solfataricus.
OC Archaea; Crenarchaeota; Thermoprotei; Sulfolobales; Sulfolobaceae;
OC Sulfolobus.
OX NCBI_TaxID=2287;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 35092 / DSM 1617 / P2;
RX MEDLINE=21332296; PubMed=11427726;
RA She Q., Singh R.K., Confalonieri F., Zivanovic Y., Allard G.,
RA Awayez M.J., Chan-Weher C.C.-Y., Clausen I.G., Curtis B.A.,
RA De Moore A., Braubo G., Fletcher C., Gordon P.M.K.,
RA Heikamp-de Jong I., Jeffries A.C., Kozera C.J., Medina N., Peng X.,
RA Thi-Ngoc H.P., Redder P., Schenk M.E., Theriault C., Tolstrup N.,
RA Charlebois R.L., Doolittle W.F., Duguet M., Gaasterland T.,
RA Garrett R.A., Ragan M.A., Sensen C.W., Van der Oost J.,
RT "The complete genome of the crenarchaeon Sulfolobus solfataricus P2.";
RL Proc. Natl. Acad. Sci. U.S.A. 98:7835-7840(2001).
DR EMBL; AE006894; AAK43138.1; -.
DR PIR; C90485; C90485.
DR HSSP; P08236; 1BHG.
DR GO; GO:0004566; F:beta-glucuronidase activity; IEA.
DR GO; GO:0005975; P:carbohydrate metabolism; IEA.
DR InterPro; IPR008979; Gal_bind_1like.
DR InterPro; IPR006101; Glyco_hydro_2ig.
DR InterPro; IPR006102; Glyco_hydro_2ig.
DR InterPro; IPR006104; Glyco_hydro_2SB.
DR InterPro; IPR006103; Glyco_hydro_2TIM.
DR Pfam; PF00703; Glyco_hydro_2; 1.
DR Pfam; PF02836; Glyco_hydro_2_C; 1.

DR Pfam; PF02837; Glyco_hydro 2 N; 1.
DR PRINTS; PR00132; GLHIDRLASE_2.
KW Complete proteome; Glycosidase; Hydrolase.
SQ SEQUENCE 570 AA; 66795 MW; DEB2FEC8050AF189 CRC64;
Query Match 33.7%; Score 1011; DB 2; Length 570;
Best Local Similarity 39.2%; Pred. No. 1.7e-58;
Matches 230; Conservative 92; Mismatches 191; Indels 74; Gaps 14;

QY 15 LNVGNLEVTSSKDRP-----IAVGSWNEQYODLCYEEGPFYKTFYVPK 60
Db 11 LOGFWKFKIDNENTGEENGWYKLGLESEDIYVPASWNEQPKWDQFSGIAMYQKDLFVSN 70
QY 61 XLSQKHIRLYFAAVNTDCEVFLNGEKVGENHIEYLPFEVDVTGKXSGENELRVVENRL 120
Db 71 DNGNRKAMWVFEAGAYITKLMINGEYGTGEGSFTQFKFPIKLKV---NEFNKIV---V 123
QY 121 KVGGFPSKVPDSCGHTVGFSGSPPPAN-----FDFPYGGIIRPVLIEFTDHARILDIM 174
Db 124 KIDNTSPSPY-----NLPPARDLNNAAFDFFNYGGIHRPVYIEFVDECHVEDIT 171
QY 175 VDTSESEPEKKLGKVKKIEVSEAVGQEMTIKLGEEKKI---RTSNRVEGEFLEN 230
Db 172 VYT-----KSYGHLKVEI-LSECNQRFSLRFLVDKEGRVILNESSNEVEKED--VNN 222
QY 231 ARFWSLEDPYLPKLELE-----KDEYTLDIGIRTI SWDEKRLYLNGKPVFLKGFGKHE 285
Db 223 VIPWSPDNPLYTLTIVEMVYGNLKDYSVERIGFRDVEVXDKIYLNKGIIFLKGFGRHE 282
QY 286 EFPVLGGTFYPLMIKDFNLKWINANSFRTSHYPYSEEWDLADRLGILVIDEAP--HV 343
Db 283 DFPILGKFTYGAVLVRDFYLMRKIGANSFRTSHYPYSENEHDLADMGFLVLEPFLCYV 342
QY 344 GITRHYNPETOKI-----AEDNIRMIDRHKHPSVIMSVANEPESNHPDAEGF 394
Db 343 NISRVMSQEEIAKMGDVXYFEKVRDTIKEMIRQHKHPSVIMSVANEPESDIREVAEF 402
QY 395 FKALYETANENDRTRPVVMVSMMDAPDERTRDVALKYPDIVCVNRYGWTYYQRIEEGL 454
Db 403 IRREVELFKSIDSSRPVTPAS-----HRSVRDLALEYVDVISLNYHGWTEWDIDISGV 457
QY 455 QALEKDIEELYARH-RKPIFVTEFGADAIAGIHYDPQPMFSEEQAEIYEKTI RLLKKD 513
Db 458 KVAIAELEIETHKKFPEKPIITTEFGADAIYGLHSDPPQMWSEEQSEMIRKYIEALREXD 517
QY 514 YIIGTHVMAFADFKTPQNVRRPILNHKGVFTDRDQPKLVAVHLRLM 560
Db 518 YIVGFHIMWVADFRTQPNSRTILNRKGI FTDRDQPKLAAKVEELF 564

RESULT 3
Q8XP19 PRELIMINARY; PRT; 599 AA.

ID Q8XP19 AC Q8XP19; PRT; 599 AA.
AC Q8XP19; PRT; 599 AA.
DT 01-MAR-2002 (TReMBLrel. 20, Created)
DT 01-MAR-2002 (TReMBLrel. 20, Last sequence update)
DT 01-OCT-2003 (TReMBLrel. 25, Last annotation update)
DE Beta-glucuronidase.
GN Name=bg1R; Ordered locusNames=CPE0147;
OS Clostridium perfringens.
OC Bacteria; Firmicutes; Clostridia; Clostridiales; Clostridiaceae;
OC Clostridium.
OX NCBI_TaxID=1502;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=13;
RX MEDLINE=21664373; PubMed=11792842;
RA Shimizu T., Ohtani K., Hirakawa H., Ohshima K., Yamashita A.,
RA Shiba T., Ogasawara N., Hattori M., Kuhara S., Hayashi H.,
RT "Complete genome sequence of Clostridium perfringens, an anaerobic
flesh-eater.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:996-1001(2002).
DR EMBL; AP003185; BAB79853.1; -.

Wed Jan 26 08:15:38 2005


```

Db      241 TTCTCAGAGAGAGAAAGTGGAGAGAAATCAATTGAATACCTTCCCTTGAAAGTAGAT 300
Qy      301 GTGACGGGGAAGTGAATCCGAGAGAAAGCACTCAGGGTGTGTTGAGAACAGATTG 360
Db      301 GTGACGGGGAAGTGAATCCGAGAGAAAGCACTCAGGGTGTGTTGAGAACAGATTG 360
Qy      361 AAAGTGGAGAGATTTCCCTCGAAAGTTCAGACAGCGGCACTCACCCGTGGATTTTT 420
Db      361 AAAGTGGAGAGATTTCCCTCGAAAGTTCAGACAGCGGCACTCACCCGTGGATTTTT 420
Qy      421 GGAAGTTTCCACCTGCAAACTTCGACTTCTCCCTACGGTGAATCATAGGCTGT 480
Db      421 GGAAGTTTCCACCTGCAAACTTCGACTTCTCCCTACGGTGAATCATAGGCTGT 480
Qy      481 CTGATAGAGTTCAAGACCAAGAGATCTGACATCTGGTGAACAGAGTGAAGTCT 540
Db      481 CTGATAGAGTTCAAGACCAAGAGATCTGACATCTGGTGAACAGAGTGAAGTCT 540
Qy      541 GAACCGGAGAGAACTTGGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGA 600
Db      541 GAACCGGAGAGAACTTGGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGA 600
Qy      601 GGACAGAGAGATGACGATCAAACTTGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 660
Db      601 GGACAGAGAGATGACGATCAAACTTGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 660
Qy      661 TTGCTCGAAGGGGAGTTCACTCTCGAAGAGCCGAGTTCTGAGCCTCGAAGATCCATAT 720
Db      661 TTGCTCGAAGGGGAGTTCACTCTCGAAGAGCCGAGTTCTGAGCCTCGAAGATCCATAT 720
Qy      721 CTTATCTCTCTCAAGTGGAACTTGAAAGAGAGAGAGAGAGAGAGAGAGAGAGAG 780
Db      721 CTTATCTCTCTCAAGTGGAACTTGAAAGAGAGAGAGAGAGAGAGAGAGAGAGAG 780
Qy      781 ACGATCAGCTGGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 840
Db      781 ACGATCAGCTGGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 840
Qy      841 TTTGGAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 900
Db      841 TTTGGAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 900
Qy      901 AAAGACTTCAACCTTCTGAAGTGAATCAACGCGAATTTCTTGAGACCTCTCACTATCT 960
Db      901 AAAGACTTCAACCTTCTGAAGTGAATCAACGCGAATTTCTTGAGACCTCTCACTATCT 960
Qy      961 TACAGTGAAGAGTGGCTGGATCTTGCCGAGAGAGAGAGAGAGAGAGAGAGAGAG 1020
Db      961 TACAGTGAAGAGTGGCTGGATCTTGCCGAGAGAGAGAGAGAGAGAGAGAGAGAG 1020
Qy      1021 CCGCAGCTTGGTATCACAAGGTACCACTACAATCCCGAGAGACTCAGAGATAGCAGAGAG 1080
Db      1021 CCGCAGCTTGGTATCACAAGGTACCACTACAATCCCGAGAGACTCAGAGATAGCAGAGAG 1080
Qy      1081 AACATAAGAGAGATATCGACAGACAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1140
Db      1081 AACATAAGAGAGATATCGACAGACAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1140
Qy      1141 GCGAAGCAACAGAGTCCATCCAGAGCGGAGGTTTCTTCAAGCCCTTTATGAG 1200
Db      1141 GCGAAGCAACAGAGTCCATCCAGAGCGGAGGTTTCTTCAAGCCCTTTATGAG 1200
Qy      1201 ACTGCCAATGAATATGATGCAACAGCGCCGCTGTGATGAGCATGATGAGAGAGAG 1260
Db      1201 ACTGCCAATGAATATGATGCAACAGCGCCGCTGTGATGAGCATGATGAGAGAGAG 1260
Qy      1261 GACGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1320
Db      1261 GACGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1320
Qy      1321 TACGGCTGTACATCTATCAGGAGAGATAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1380
Db      1321 TACGGCTGTACATCTATCAGGAGAGATAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1380

```

```

Db      1321 TACGGCTGTACATCTATCAGGAGAGATAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1380
Qy      1381 ATAGAGAGCTTATGCAAGGACAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1440
Db      1381 ATAGAGAGCTTATGCAAGGACAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1440
Qy      1441 GCGATAGCTGGCATCCACTACGATCCAGTCCAAATGTTCTCCGAGAGATACCAAGCAGAG 1500
Db      1441 GCGATAGCTGGCATCCACTACGATCCAGTCCAAATGTTCTCCGAGAGATACCAAGCAGAG 1500
Qy      1501 CTCGTTGAAAGAGAGATGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1560
Db      1501 CTCGTTGAAAGAGAGATGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1560
Qy      1561 TGGGCTTTGAGAGATTTTAAGACTCCTCAGAGATGAGAGAGAGAGAGAGAGAGAGAGAG 1620
Db      1561 TGGGCTTTGAGAGATTTTAAGACTCCTCAGAGATGAGAGAGAGAGAGAGAGAGAGAGAG 1620
Qy      1621 GGTGTTTTCACAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1680
Db      1621 GGTGTTTTCACAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1680
Qy      1681 AGTGAGGTT 1689
Db      1681 AGTGAGGTT 1689

```

```

RESULT 2
AE001766/c 12583 bp DNA linear BCT 04-JUN-2004
LOCUS Thermotoga maritima MSB section 78 of 136 of the complete genome.
DEFINITION AE001766 AE000512
ACCESSION AE001766.1 GI:4981600
KEYWORDS
SOURCE Thermotoga maritima MSB
ORGANISM Thermotoga maritima MSB
REFERENCE 1 (bases 1 to 12583)
AUTHORS Nelson,K.E., Clayton,R.A., Gill,S.R., Gwinn,M.L., Dodson,R.J.,
Hart,D.H., Hickey,E.K., Peterson,J.D., Nelson,W.C., Ketchum,K.A.,
McDonald,L., Utterback,T.R., Malek,J.A., Linher,K.D., Garrett,M.M.,
Stewart,A.M., Cotton,M.D., Pratt,M.S., Phillips,C.A.,
Richardson,D., Heidelberg,J., Sutton,G.G., Fleischmann,R.D.,
Eisen,J.A., White,O., Salzberg,S.L., Smith,H.O., Venter,J.C. and
Fraser,C.M.
Evidene for lateral gene transfer between Archaea and bacteria
from genome sequence of Thermotoga maritima
Nature 399 (6734), 323-329 (1999)

```

```

TITLE
JOURNAL
MEDLINE
PUBMED
REFERENCE
AUTHORS
2 (bases 1 to 12583)
Nelson,K.E., Clayton,R.A., Gill,S.R., Gwinn,M.L., Dodson,R.J.,
Hart,D.H., Hickey,E.K., Peterson,J.D., Nelson,W.C., Ketchum,K.A.,
McDonald,L., Utterback,T.R., Malek,J.A., Linher,K.D., Garrett,M.M.,
Stewart,A.M., Cotton,M.D., Pratt,M.S., Phillips,C.A.,
Richardson,D., Heidelberg,J., Sutton,G.G., Fleischmann,R.D.,
White,O., Salzberg,S.L., Smith,H.O., Venter,J.C. and Fraser,C.M.
Direct Submission
Submitted (01-JUN-1999) The Institute for Genomic Research, 9712
Medical Center Dr, Rockville, MD 20850, USA

```

```

FEATURES
Source
1.12583
/organism="Thermotoga maritima MSB"
/mol_type="genomic DNA"
/strain="MSB"
/db_xref="taxon:243274"
81.1607
/locus_tag="TM1058"
81.1607
/locus_tag="TM1058"

```

```

CDs
/locus_tag="TM1058"
/locus_tag="TM1058"
/note="similar to GB:AE000666 percent identity: 48.70;
identified by sequence similarity; putative"
/codon_start=1

```

```

/transl_table=11
/product="glutamate synthase-related protein"
/protein_id="AAD36135.1"
/db_xref="GI:4981601"
/translation="MSGKCAKCDGCTGCTETWLASFRGVEVYGPFGDITAGAVKD
VVDYSHNLNIGYARGAEGLEVEPPDPTAFTNVDTTTEYMDIKVMKVDIFETGA
LGSTEIARKMDHIAVGAIGITVCGENVAGVDPDLDSNGKVKSPELDRIE
YKRYHDEYGEILLIOMNVEDTRGVAEYVINKGIEITELKMGAKSIGELIKVRS
BRALELRKGYVLPDPPEVORAFKEGEIKEPERHSLGFVSKESFLKEVERLKL
GPKRITLKTGAYSAVELAMALRYGAEKVDLIVDGA PGGTGMSPMNENWGIPTFY
LEALTYQFAEKLRRGIRVDPDIALAGSESTEDVLKAIAMGSPYKAVCMGRALMI PA
MVGNIIGEWLKSGLPKTVSKYGTVEIIVTYEELRSRFGEEYKPLGAIIVYTFE
VQFKRTGLQQLMAGARKFRSLSRKDLIALTKDAEISGIPYVESYRDEAERLLEE
"
complement(1650..1913)
/locus_tag="TM1059"
complement(1650..1913)
/locus_tag="TM1059"
/notes="similar to SP:P45693 PID:862985 GB:AL009126 percent
identity: 48.70; identified by sequence similarity;
putative"
/codon_start=1
/transl_table=11
/product="spovs-related protein"
/protein_id="AAD36136.1"
/db_xref="GI:4981602"
/translation="MEVLKVSXSKSDPNKAGAIAGVREHGKAEIQAGAVNQAVK
AIAIRGYLAPSGIDLVFPAFTVEIENKRTAIFVPPKS"
complement(1962..3272)
/locus_tag="TM1060"
complement(1962..3272)
/locus_tag="TM1060"
/notes="similar to GB:AE00782 percent identity: 49.74;
identified by sequence similarity; putative"
/codon_start=1
/transl_table=11
/product="conserved hypothetical protein"
/protein_id="AAD36144.1"
/db_xref="GI:4981610"
/translation="MNISIEGALAVLINQFEGAYLTGYFLMWGASSFFIGLFGSIP
FLANTLQLTLSFSHRLKSRKQIIVPLMTARTSILLFAVPAIKHGLLAYLYFYI
FIAGALSAPLMQSMWSDLPKDMGYSFGFRNLHGAVQIPAMELAGAILDSLENMK
GFGTLFIAGSLGALNGYFLKIQYEPYKPREASVITKAVKFLKEHFKNLFJGFA
FMNFAIGVGVYINVMILKEVESYLOISVLNAGMFIQTLFQDFWGLGDRYGFQYF
LKVCLMHAIVILMTLTPRSFLVFLQIITIGFTAGTSQLVFYTLMTAPSSLKT
EAFSVNSLSNLSIFAGSLVAVSLNISLPGISAIRLTMFISFLRASAAVYII
SRMDLGTPOKVDSLIQAVKESFSGTVPWIRERLNTLNIFFRRKR"
complement(3382..4545)
/locus_tag="TM1061"
complement(3382..4545)
/locus_tag="TM1061"
/notes="similar to percent identity: 0.00; identified by
sequence similarity; putative"
/codon_start=1
/transl_table=11
/product="hypothetical protein"
/protein_id="AAD36137.1"
/db_xref="GI:4981603"
/translation="MRSMTDVRRYWISLRLKICEMPLECASDRLLKSMPEVGKSEE
RRKFTLLELGRIFCGISPLELNRESPTDPEERKIAIRLSEFAVKSIDVATPNCK
DYNMFKGRQPLVDAAFLVEAIIARPKVWEDLVSTKRLIRLKAIRKIEPYPSNM
LIFSAMETFEFFPAGEWSDTKVDLILKNVESWYKGDGAPFFRMDYNSFVIYP
MTIDVLRITISEKTEWKELYVKVLRARQYAVLERMISPEGTPIIGRSITRTAVF
HLISQLSLHLPLASLSPAVRCALTAVLRIFENSTPDENGMLKIGVIGSQPSLGE
EYITGSLVCTVFLPLGPTSDPFWRDPCKKTKKWEGEDVAPDRALIED"
complement(4542..6233)
/locus_tag="TM1062"
complement(4542..6233)
/locus_tag="TM1062"
/notes="similar to PID:642973 PID:14446 PID:412358 percent
identity: 57.73; identified by sequence similarity;
putative"
/codon_start=1

```

```

/transl_table=11
/product="beta-glucuronidase"
/protein_id="AAD36143.1"
/db_xref="GI:4981609"
/translation="MVRPQRNKKRFTILLNGVWNLVETSKDRPIAVPGSMNEOYDLC
YBGFPTVKTTFYVPEKLSQKHILYPAVNTDCEVFLGKEKGVENHIEYLPFEVDVT
GKVKSGENELRVVENRLKVGGFPSKVPDSTHTVGFPSFPANFDPFGGIIIPV
LIEFTDARILDIWVDTSESEPEKLGKVKKIEVSEAVQEMTILKGBEEKIRTS
NRFVEGEFLLENARLSDLPDYLPLKLEKDEYTLIDGIRTISDEKRLYNGKPV
FLKFGKHFEFPVLGGTFPDLIKDFNLKWINANSFRTSHYPYSEEWLADRLGI
LVIDEAPHVGIITRYHNPETQKIAEDNIRKMDIRKKNPSVIMSVANEPESNDPAE
GFPKALYETANEMDRTRPVVMQMDAPDERTDVALKYFDIIVCNRYGWIYQGR
BEGLOALEXDIIEELYARHKKPIFTEFGADAIAGIHYPDQMESEYQAEIVKTI RL
LLKDYIIGTHWAFADFKTPQNVRRPILNKGVTFRDROPKLVAVHRLMSEV"
complement(6284..7309)
/locus_tag="TM1063"
complement(6284..7309)
/locus_tag="TM1063"
/notes="similar to SP:P42065 PID:677944 GB:AL009126 percent
identity: 75.31; identified by sequence similarity;
putative"
/codon_start=1
/transl_table=11
/product="oligopeptide ABC transporter, ATP-binding
protein"
/protein_id="AAD36138.1"
/db_xref="GI:4981604"
/translation="WALLEVKNLKKYEPILIKRGFRKRVGVLKAVDGISFHIIEGTL
GLVSGCGCTTTAKLIMRGEIPEGEIVLNMDGEKVDITKLEKELREKGVRFLOM
IFQDPYSSLNPRMTARDIIAEPVNVGVKGEIDAVSDILRAVGRPEYMOYRPH
AFSGGQRORIAIARAIALPKPLVLCDEPTALDVSVQAOIINLKDLOKEYNLTYLFI
SHDLGVVEHITNRVAVMYVGRIVELAEETELFSSPKHPTALSLAVPKDPKRRK
AQLTGEVDPDSNPSPGCVFHPRCFYAKAIKCEVYPLDRNVGTQDNPLVACHFADSLK
LEGVGTM"
complement(7296..8294)
/locus_tag="TM1064"
complement(7296..8294)
/locus_tag="TM1064"
/notes="similar to SP:P42064 PID:677943 GB:AL009126 percent
identity: 72.95; identified by sequence similarity;
putative"
/codon_start=1
/transl_table=11
/product="oligopeptide ABC transporter, ATP-binding
protein"
/protein_id="AAD36139.1"
/db_xref="GI:4981605"
/translation="MSTLLQIKNLRTYFTDEGVKAVDGVSEIEEGRTLGVGESG
CGKSVTARSIIKLSTAGRIYSGEILYNDQMDQVLDVKSKEIRKVRGRHIAIMFOE
PMASFSPVITIGDQITEGMVYHFGITKQEARERAVELLRRVGIKPKEMIDSYPEYS
GMAFORAMIMALSCNPRLLADEPTTALDVYIOAQVLDLKDLOQOEYKMAIMITHN
MGVAEMADHVVMYLGKRVESAPVEELFYNDKHPYTSLSIRSIPIVGRKVERLEVI
GDVPDPNMPKGRFHPRCFYMKWICIDERPEVEVEGPEHRVSCFLYGEKDGAS"
complement(8306..9409)
/locus_tag="TM1065"
complement(8306..9409)
/locus_tag="TM1065"
/notes="similar to GB:AE000657 percent identity: 67.80;
identified by sequence similarity; putative"
/codon_start=1
/transl_table=11
/product="oligopeptide ABC transporter, permease protein"
/protein_id="AAD36140.1"
/db_xref="GI:4981606"
/translation="MKKREQMEKEFYASOMOLIMWRFRKHLAVIGVLLIIVYFA
IFCEFPAPYDPNKNYRYPYAPORIHFEHGKFGIPFVGYTYVDLETLRIYKED
KSKIKFKIFVRGDEYKFWGIWKTNIHIGVEDGHMFLGTDLSGRMLSRITYGARI
STSIGLIGVFLSFLIGVAIGISGYFGCAVDNFIOQTIEIKSIPITPLMLASALP
QNMPLRVYFVIVILSLTGWTDLARYRSRLSLREDPFVMAAKFGASGARIIFRH
MLPSFMSHLIASITLSIPGMITGETSLSLGLGRPVIISWGVLLQEAQNLTVVALYP
WLLIPVVFVITVLCFNFVGDGLRDADPYANM"
complement(9420..10403)
/locus_tag="TM1066"

```

CDS		complement(9420..10403)	
		/locus_tag="TM106"	
		/note="similar to SP:P42062 PID:677946 GB:AL009126 percent identity: 62.93; identified by sequence similarity; putative"	
		/codon_start=1	
		/transl_table=11	
		/product="oligopeptide ABC transporter, permease protein"	
		/protein_id="AAD36141.1"	
		/db_xref="GI:4981607"	
Query Match		99.9%;	Score 1688; DB 1; Length 12583;
Best Local Similarity		99.9%;	Pred. No. 0;
Matches 1688;		Conservative 0;	Mismatches 1; Indels 0; Gaps 0;
QY	1	ATGTAAGACCGCAACGAAACAAGAAGATTATTTCTTATCTTGAATGAGTTGGAAT	60
DB	6233	ATGTAAGACCGCAACGAAACAAGAAGATTATTTCTTATCTTGAATGAGTTGGAAT	6174
QY	61	CTTGAAGTAACCAAGACAGACCAATCGCCGTTCTGGAAGCTGGAATGAGCAGTAC	120
DB	6173	CTTGAAGTAACCAAGACAGACCAATCGCCGTTCTGGAAGCTGGAATGAGCAGTAC	6114
QY	121	CAGATCTGTGTACGAAGAAGACCCCTTCACTTCAAAAACCACTTCTACGTTCCGAAG	180
DB	6113	CAGATCTGTGTACGAAGAAGACCCCTTCACTTCAAAAACCACTTCTACGTTCCGAAG	6054
QY	181	NAACCTTCACAAAAAACATCAGACTTTACTTGTGCGGTGAACAACGAGCTGCCAGTTC	240
DB	6053	GAACCTTCACAAAAAACATCAGACTTTACTTGTGCGGTGAACAACGAGCTGCCAGTTC	5994
QY	241	TTCTTCAACGAGAGAAAGTGGAGAAATCACAATTGAATACCTTCCCTTGAAGTAGAT	300
DB	5993	TTCTTCAACGAGAGAAAGTGGAGAAATCACAATTGAATACCTTCCCTTGAAGTAGAT	5934
QY	301	GTAACGGGGAAAGTGAATCCGAGAGAAACGAATCAGGGGTGTTGAGAACAGATTG	360
DB	5933	GTAACGGGGAAAGTGAATCCGAGAGAAACGAATCAGGGGTGTTGAGAACAGATTG	5874
QY	361	AAAGTGGAGAGATTTCCTCGAAGTTCCAGACAGCGGCACTCACACCGTGGATTTTTT	420
DB	5873	AAAGTGGAGAGATTTCCTCGAAGTTCCAGACAGCGGCACTCACACCGTGGATTTTTT	5814
QY	421	GGAAGTTTTCACCTGCAAACTTTCGACTTCTTCCCTACGGTGAATCATAGGCTGTT	480
DB	5813	GGAAGTTTTCACCTGCAAACTTTCGACTTCTTCCCTACGGTGAATCATAGGCTGTT	5754
QY	481	CTGATAGAGTTCAACACCAACGCGAGATCTCGACATCTGGGTGAACACGAGTAGTCT	540
DB	5753	CTGATAGAGTTCAACACCAACGCGAGATCTCGACATCTGGGTGAACACGAGTAGTCT	5694
QY	541	GAAACCGGAGAACTTGGAAAAGTGAAGATGAAGTCTCAGAAAGAACGGGTG	600
DB	5693	GAAACCGGAGAACTTGGAAAAGTGAAGATGAAGTCTCAGAAAGAACGGGTG	5634
QY	601	GGACAGAGAGATGACGATCAAACTTGGAGAGAGAGAAAAAGATTAGAACATCCAAACGA	660
DB	5633	GGACAGAGAGATGACGATCAAACTTGGAGAGAGAGAAAAAGATTAGAACATCCAAACGA	5574
QY	661	TTTGTGGAAGGGAGTTTCTCTGAAAAACGCCAGGTTCTGGAGCTCGAAGATCCATAT	720
DB	5573	TTTGTGGAAGGGAGTTTCTCTGAAAAACGCCAGGTTCTGGAGCTCGAAGATCCATAT	5514
QY	721	CTTTATCTCTCAAGGTGAACCTTGAAGAGAGAGTACACTCTGGAACATCGGAATCAGA	780
DB	5513	CTTTATCTCTCTCAAGGTGAACCTTGAAGAGAGAGTACACTCTGGAACATCGGAATCAGA	5454
QY	781	ACGATCAGCTGGAGAGAGAGAGGCTTATCTGAACGGGAAACCTGCTTTTGAAGGGC	840
DB	5453	ACGATCAGCTGGAGAGAGAGAGGCTTATCTGAACGGGAAACCTGCTTTTGAAGGGC	5394
QY	841	TTTGGAAAGCAGAGGAATTCCCGGTTCTGGGGCAGGGCACCTTTTATCCATTGATGATA	900

DB	5393	TTTGGAAAGCAGAGGAATTCCCGGTTCTGGGGCAGGGCACCTTTTATCCATTGATGATA	5334
QY	901	AAAGACTTCAACCTTCTGAAGTGAATCAACGGCAATTCTTTGAGAACCTCTCACTATCCT	960
DB	5333	AAAGACTTCAACCTTCTGAAGTGAATCAACGGCAATTCTTTGAGAACCTCTCACTATCCT	5274
QY	961	TACAGTGAAGAGTGGCTGATCTTGCCGACAGACTCGGAATCCTTGATAGAGAAACC	1020
DB	5273	TACAGTGAAGAGTGGCTGATCTTGCCGACAGACTCGGAATCCTTGATAGAGAAACC	5214
QY	1021	CCGACGTTGGTATCACAAGGTACCACTACAAATCCCGAGACTCAGAAGATAGCAAGAAC	1080
DB	5213	CCGACGTTGGTATCACAAGGTACCACTACAAATCCCGAGACTCAGAAGATAGCAAGAAC	5154
QY	1081	AACATAAGAAAGATGATCGACAGACACAAGAACCATCCAGTGTGATCATGTGAGTGTG	1140
DB	5153	AACATAAGAAAGATGATCGACAGACACAAGAACCATCCAGTGTGATCATGTGAGTGTG	5094
QY	1141	GCGAACGAACCAAGTCCAAACCATCCAGACGGGAGGGTTCTTCAAAAGCCCTTTATGAG	1200
DB	5093	GCGAACGAACCAAGTCCAAACCATCCAGACGGGAGGGTTCTTCAAAAGCCCTTTATGAG	5034
QY	1201	ACTGCCAATGAATGATCGAACACAGCCCGTTGTCTATGTTGATGATGATGACGACCA	1260
DB	5033	ACTGCCAATGAATGATCGAACACAGCCCGTTGTCTATGTTGATGATGATGACGACCA	4974
QY	1261	GACGAGAGAAAGAGACGTGGCGCTGAAGTACTTGCACATGCTGTGTGAACAGGTAC	1320
DB	4973	GACGAGAGAAAGAGACGTGGCGCTGAAGTACTTGCACATGCTGTGTGAACAGGTAC	4914
QY	1321	TACGGCTGTACATCTATCAGGGAAAGATAGAAAGAGACTTCAAGCTCTGAAAAAGAC	1380
DB	4913	TACGGCTGTACATCTATCAGGGAAAGATAGAAAGAGACTTCAAGCTCTGAAAAAGAC	4854
QY	1381	ATAGAAGAGCTTATGCAAGGCAAGAAAGCCATCTTGTCAAGAAATTCGGTCCGAC	1440
DB	4853	ATAGAAGAGCTTATGCAAGGCAAGAAAGCCATCTTGTCAAGAAATTCGGTCCGAC	4794
QY	1441	GCGATAGCTGGCATCCACTACGATCCACCTCAATGTTCTCCGAAGAGTACCAACAGAG	1500
DB	4793	GCGATAGCTGGCATCCACTACGATCCACCTCAATGTTCTCCGAAGAGTACCAACAGAG	4734
QY	1501	CTCGTTGAAAAGACGATCAGGCTCTTTGAAAAAAGACTACATCATCGGAACACAGTG	1560
DB	4733	CTCGTTGAAAAGACGATCAGGCTCTTTGAAAAAAGACTACATCATCGGAACACAGTG	4674
QY	1561	TGGGCTTTGAGATTTTAAGACTCTCAGAAATGTGAGAAAGCCATTCTCAACACAAG	1620
DB	4673	TGGGCTTTGAGATTTTAAGACTCTCAGAAATGTGAGAAAGCCATTCTCAACACAAG	4614
QY	1621	GGTGTTTTCAAGAGACAGACAACCAACTCGTGTCTATGTACTGAGAAGACTGTGG	1680
DB	4613	GGTGTTTTCAAGAGACAGACAACCAACTCGTGTCTATGTACTGAGAAGACTGTGG	4554
QY	1681	AGTGAGGTT 1689	
DB	4553	AGTGAGGTT 4545	

RESULT 3	AF012423	2153 bp	mRNA	linear	MAM 09-SEP-1999
LOCUS	AF012423				
DEFINITION	Felis catus beta-glucuronidase (GUSB) mRNA, complete cds.				
ACCESSION	AF012423				
VERSION	AF012423.1	GI:4102550			
KEYWORDS					
SOURCE	Felis catus (cat)				
ORGANISM	Felis catus				
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Carnivora; Fissipedia; Felidae; Felis.				
AUTHORS	Fyfe,J.C., Kurzhals,R.L., Lassaline,M.E., Henthorn,P.S., Alur,P.R., Wang,P., Wolfe,J.H., Giger,U., Haskins,M.E., Patterson,D.F.,				

